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AMENDMENTS TO THE CLAIMS

1. (Currently amended) A pharmaceutical composition for treating or preventing an allergic reaction associated with increased IgE levels or inhibiting cellular proliferation in a mammal comprising anycompound or salt thereof represented by one or more of the following compounds formulas:

wherein Q, T, X, and Z are independently selected from N or C, and wherein one of Q, T, X, and Z is N;

wherein A is selected from the group consisting of H, halogen, and CONHR₁; wherein n is a number from one to four;

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wherein L and M are independently selected from the group consisting of H, alkyl, alkoxy, aryl, substituted aryl, hydroxy, halogen, amino, alkylamino, nitro, cyano, CF₃, OCF₃, CONH₂, CONHR and NHCOR₁;

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur.

2. (Currently amended) A-pharmaceutical composition The compound or salt thereof of Claim 1, wherein the Supragenus A-D are represented by the formulas:

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Genus C4; Genus D1; Genus D2; Genus D3; and

wherein Q, T, X, and Z are independently selected from N or C, and wherein one of Q, T, X, and Z is N;

Genus D4;

wherein A is selected from the group consisting of H and halogen;

wherein n is a number from one to four;

wherein L and M are independently selected from the group consisting of H, alkyl, alkoxy, aryl, substituted aryl, hydroxy, halogen, amino, alkylamino, nitro, cyano, CF₃, OCF₃, CONH₂, CONHR and NHCOR₁;

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wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur.

- 3. (Currently amended) The compound <u>or salt thereof</u> of Claim 1, wherein said polycyclic aliphatic group is selected from the group consisting of adamantyl, bicycloheptyl, camphoryl, bicyclo[2,2,2]octanyl and norbornyl.
- 4. (Currently amended) The compound <u>or salt thereof</u> of Claim 1, wherein said heteroaryl and said substituted heteroaryl is selected from the group consisting of pyridines, thiazoles, isothiazoles, oxazoles, pyrimidines, pyrazines, furans, thiophenes, isoxazoles, pyrroles, pyridazines, 1,2,3-triazines, 1,2,4-triazines, 1,3,5-triazines, pyrazoles, imidazoles, indoles, quinolines, iso-quinolines, benzothiophines, benzofurans, parathiazines, pyrans[[,]] <u>and</u>

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chromenes, pyrrolidines, pyrazolidines, imidazolidines, morpholines, thiomorpholines, and the corresponding heterocyclics.

- 5. (Cancelled)
- 6. (Currently amended) The <u>compound or salt thereof pharmaceutical composition</u> of Claim 1, wherein R₁ and R₂ are independently selected from the group consisting of:

- 7. (Currently amended) The pharmaceutical composition compound or salt thereof of Claim 1-comprising a compound-selected from the group consisting of compounds S-1 to S-25.
- 8. (Currently amended) A pharmaceutical composition for treating or preventing an allergic reaction associated with increased IgE levels or inhibiting cellular proliferation in a mammal comprising anycompound or salt thereof represented by one or more of the following compounds formula:

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wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur.

- 9. (Currently amended) The pharmaceutical composition compound or salt thereof of Claim 8-comprising a compound selected from the group consisting of compounds S-4, S-5, S-6, S-7, S-8, S-11, S-13, S-15 and S-16.
- 10. (Currently amended) The pharmaceutical composition compound or salt thereof of Claim 9 comprising represented by the compound S-7.

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11. (Currently amended) A <u>compound or salt thereof represented by pharmaceutical</u> composition for treating or preventing an allergic reaction associated with increased IgE levels or inhibiting cellular proliferation in a mammal comprising any one or more of the following compounds formula:

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur.

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12. (Currently amended) The <u>compound or salt thereof pharmaceutical composition of</u> Claim 11 comprising a compound selected from the group consisting of compounds S-17, S-19, S-20[[,]] and S-21.

13. (Currently amended) A <u>compound or salt thereof represented by pharmaceutical</u> composition for treating or preventing an allergic reaction associated with increased IgE levels or inhibiting cellular proliferation in a mammal comprising any one or more of the following compounds formula:

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl

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contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur.

- 14. (Currently amended) The <u>compound or salt thereof pharmaceutical composition of</u> Claim 13 <u>comprising represented by the compound S-24.</u>
- 15. (Currently amended) A <u>compound or salt thereof represented by pharmaceutical</u> composition for treating or preventing an allergic reaction associated with increased IgE levels or inhibiting cellular proliferation in a mammal comprising any one or more of the following compounds formula:

$$\begin{array}{c|c} R_1 - NH & & & \\ & & & \\ & & & \\ R_1 - N & & \\ & & &$$

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

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wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur.

- 16. (Currently amended) The <u>compound or salt thereof pharmaceutical composition of</u> Claim 15 <u>comprising represented by the compound S-25.</u>
- 17. (Currently amended) A <u>compound or salt thereof represented by pharmaceutical</u> composition for treating or preventing an allergic reaction associated with increased IgE levels or inhibiting cellular proliferation in a mammal comprising any one or more of the following compounds formula:

$$(A)_{n} \xrightarrow{T} \underbrace{N}_{N} \underbrace{N}_{$$

wherein T and X are independently selected from N or C, and wherein one of T and X is N;

wherein A is selected from the group consisting of H, halogen, and CONHR₁; wherein n is a number from one to four;

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

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wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur.

18. (Withdrawn – currently amended) A method for treating or preventing an allergic reaction and/or for inhibiting cytokines or leukocytes in a mammal comprising administering an effective amount of any one or more of the following compounds or salts thereof of Claim 1.[[:]]

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wherein Q, T, X, and Z are independently selected from N or C, and wherein one of Q, T, X, and Z is N;

wherein A is selected from the group consisting of H, halogen, and CONHR₁; wherein n is a number from one to four;

wherein L and M are independently selected from the group consisting of H, alkyl, alkoxy, aryl, substituted aryl, hydroxy, halogen, amino, alkylamino, nitro, cyano, CF₃, OCF₃, CONH₂, CONHR and NHCOR₁;

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ eyeloalkyl, substituted C₃-C₉ eyeloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and

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substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur.

- 19. (Withdrawn) The method of Claim 18 further comprising administering at least one additional ingredient which is active in reducing at least one symptom associated with said allergic reaction.
- 20. (Withdrawn) The method of Claim 19, wherein said at least one additional ingredient is selected from the group consisting of a short-acting β₂-adrenergic agonist, a longacting \(\beta_2\)-adrenergic agonist, an antihistamine, a phosphodiesterase inhibitor, an anticholinergic agent, a corticosteroid, an inflammatory mediator release inhibitor and a leukotriene receptor antagonist.
- 21. (Withdrawn) The method of Claim 19, wherein said at least one additional ingredient is combined with said compound in a pharmaceutically acceptable diluent and coadministered to the mammal.
- 22. (Withdrawn) The method of Claim 18, wherein said compound is administered at a dose of about 0.01 mg to about 100 mg per kg body weight per day.
- 23. (Withdrawn) The method of Claim 22, wherein said dose is administered in divided doses at regular periodic intervals.
- 24. (Withdrawn) The method of Claim 23, wherein said regular periodic intervals occur daily.
- 25. (Withdrawn – currently amended) A method for treating or preventing asthma in a mammal comprising administering an IgE-suppressing amount of any one or more of the following-compounds or salt thereof of Claim 1.[[:]]

$$(A)_{n} \xrightarrow{T} Q \qquad N \qquad N \qquad N \qquad N \qquad R_{2}$$

$$X \nearrow Z \qquad N \qquad M \qquad O$$

Supragenus A;

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wherein Q, T, X, and Z are independently selected from N or C, and wherein one of Q, T, X, and Z is N;

wherein A is selected from the group consisting of H, halogen, and CONHR₁; wherein n is a number from one to four;

wherein L and M are independently selected from the group consisting of H, alkyl, alkoxy, aryl, substituted aryl, hydroxy, halogen, amino, alkylamino, nitro, cyano, CF₃, CONH₂, CONHR and NHCOR₁;

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

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wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur.

- 26. (Withdrawn) The method of Claim 25 further comprising administering at least one additional ingredient which is active in reducing at least one symptom associated with said asthma.
- 27. (Withdrawn) The method of Claim 26, wherein said additional ingredient is selected from the group consisting of a short-acting β_2 -adrenergic agonist, a long-acting β_2 -adrenergic agonist, an antihistamine, a phosphodiesterase inhibitor, an anticholinergic agent, a corticosteroid, an inflammatory mediator release inhibitor and a leukotriene receptor antagonist.
- 28. (Withdrawn currently amended) A method for inhibiting cellular proliferation in a mammal comprising administering an <u>effective</u> amount of any one or more of the <u>following</u> compounds or salts thereof of Claim 1[[:]]

$$(A)_n \xrightarrow{T}^Q \underset{L}{N} \underset{N}{N} \xrightarrow{R_2}$$

Supragenus A;

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wherein Q, T, X, and Z are independently selected from N or C, and wherein one of Q, T, X, and Z is N;

wherein A is selected from the group consisting of H, halogen, and CONHR₁; wherein n is a number from one to four;

wherein L and M are independently selected from the group consisting of H, alkyl, alkoxy, aryl, substituted aryl, hydroxy, halogen, amino, alkylamino, nitro, cyano, CF₃, OCF₃, CONH₂, CONHR and NHCOR₁;

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

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wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur.

- 29. (Withdrawn) The method of Claim 28 further comprising administering at least one additional ingredient which is active in reducing at least one symptom associated with said cellular proliferation.
- 30. (Withdrawn) The method of Claim 29, wherein said at least one additional ingredient is selected from the group consisting of antifungals, antivirals, antibiotics, anti-inflammatories, and anticancer agents.
- 31. (Withdrawn) The method of Claim 29, wherein said at least one additional ingredient is selected from the group consisting of alkylating agent, antimetabolite, DNA cutter, topoisomerase I poison, topoisomerase II poison, DNA binder, and spindle poison.
- 32. (Withdrawn) The method of Claim 29, wherein said at least one additional ingredient is combined with said compound in a pharmaceutically acceptable diluent and co-administered to the mammal.
- 33. (Withdrawn) The method of Claim 28, wherein said compound is administered at a dose of about 0.01 mg to about 100 mg per kg body weight per day.
- 34. (Withdrawn) The method of Claim 33, wherein said dose is administered in divided doses at regular periodic intervals.
- 35. (Withdrawn) The method of Claim 34, wherein said regular periodic intervals occur daily.

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36. (Withdrawn) The method of Claim 28 further comprising administering at least one other therapy which is effective in ameliorating at least one symptom associated with cellular hyperproliferation.

- 37. (Withdrawn) The method of Claim 36, wherein said therapy is an anti-cancer therapy.
- 38. (Withdrawn) The method of Claim 36, wherein said therapy is selected from the group consisting of radiation, immunotherapy, gene therapy, and surgery.
- 39. (Withdrawn currently amended) A method of preparing a compound or salt thereof having the formula of Supragenus A as defined in Claim 1 comprising:

$$(A)_{n} \xrightarrow{T}^{Q} \xrightarrow{N} \xrightarrow{N} \xrightarrow{N} \xrightarrow{R_{2}}$$

Supragenus A;

wherein Q, T, X, and Z are independently selected from N or C, and wherein one of Q, T, X, and Z is N;

wherein A is selected from the group consisting of H, halogen, and CONHR₁; wherein n is a number from one to four;

wherein L and M are independently selected from the group consisting of H, alkyl, alkoxy, aryl, substituted aryl, hydroxy, halogen, amino, alkylamino, nitro, cyano, CF₃, OCF₃, CONH₂, CONHR and NHCOR₁;

wherein R is selected from the group consisting of H, C₁-C₅-alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅-alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

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wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ eycloalkyl, substituted C₃-C₉ eycloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said method comprises steps:

 $(A)_n + X_n = 0$

reacting a compound with a formula: XY.Z CI with ammonium hydroxide

 $(A)_{n} \xrightarrow{T} Q NO_{2}$ NH_{2}

to form a compound with a formula:

(A)_n; NH

reacting the compound with a formula: L' with diammonium sulfide

to form a compound with a formula:

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reacting the compound with a formula: $(A)_n \overset{\mathsf{T}}{\overset{\mathsf{Q}}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}{\overset{\mathsf{NH}_2}{\overset{\mathsf{NH}_2}{\overset{\mathsf{NH}_2}{\overset{\mathsf{NH}_2}{\overset{\mathsf{NH}_2}{\overset{\mathsf{NH}_2}{\overset{\mathsf{NH}_2}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}{\overset{\mathsf{NH}_2}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}{\overset{\mathsf{NH}_2}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{N}}}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}}}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}}{\overset{\mathsf{NH}_2}}}}}}}}}}}}}}}}}$

with a compound with a

CI NO₂

to form a compound with a formula:

 $(A)_{n} \xrightarrow{T} Q \xrightarrow{NH} O \xrightarrow{NO_{2}} M$

formula:

cyclizing the compound with a formula:

with use of

 $(A)_{n} \xrightarrow{T} Q \qquad N \qquad \qquad \downarrow$

an acid to form a compound with a formula:

 $(A)_{n} \xrightarrow{T} Q \qquad N \qquad NO_{2}$ $X \qquad Z \qquad N \qquad M$

reducing the compound with a formula:

to form a

with an acyl

compound with a formula: $(A)_{n} \xrightarrow{T}^{Q} X_{H} \xrightarrow{N}^{N} X_{H} \xrightarrow{N}^{N} X_{H}$; and

 $(A)_{n} \xrightarrow{X}_{L} Z \xrightarrow{N}_{H} M^{NH_{2}}$

reacting the compound with a formula: chloride to form a compound of Supragenus A.

40. (Withdrawn – currently amended) A method of preparing a compound or salt thereof having the formula of Supragenus A as defined in Claim 1 comprising:

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$$(A)_{n} \xrightarrow{T}^{Q} \underset{L}{\stackrel{N}{\bigvee}}_{Z} \xrightarrow{N} \underset{M}{\stackrel{H}{\bigvee}}_{Q} \xrightarrow{R_{2}}$$

Supragenus A;

wherein Q, T, X, and Z are independently selected from N or C, and wherein one of Q, T, X, and Z is N;

wherein A is selected from the group consisting of H, halogen, and CONHR₁; wherein n is a number from one to four;

wherein L and M are independently selected from the group consisting of H, alkyl, alkoxy, aryl, substituted aryl, hydroxy, halogen, amino, alkylamino, nitro, cyano, CF₃, OCF₃, CONH₂, CONHR and NHCOR₁;

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ eyeloalkyl, substituted C₃-C₉ eyeloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl

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contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said method comprises steps:

reacting a compound with a formula:

with ammonium hydroxide

to form a compound with a formula: $(A)_n \xrightarrow{T}_{Z} NH_2$

to form a compound with a formula: $(A)_{n} \stackrel{T}{\overset{Q}{\overset{\vee}{\underset{}{\bigvee}}}} Z \stackrel{NH_2}{\overset{NH_2}{\underset{}{\bigvee}}}$

reacting the compound with a formula: $(A)_n \stackrel{T}{\downarrow}_{\downarrow}^{NH_2}$ with a compound with a

H NO₂

to form a compound with a formula:

 $(A)_{n} \underbrace{\overset{T}{\overset{Q}{\overset{}}{\overset{}}{\overset{}}}}_{Z} \overset{Q}{\overset{}{\overset{}}{\overset{}}}_{H}} \overset{N}{\overset{}} \overset{NO_{2}}{\overset{}}$

formula:

reducing the compound with a formula:

 $(A)_{n} \xrightarrow{T} Q N NO_{2}$ N M to form a

with an acyl

reacting the compound with a formula: L' H M chloride to form a compound of Supragenus A.

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41. (Withdrawn – currently amended) A method of preparing a compound or salt thereof having the formula of Supragenus B as defined in Claim 1 comprising:

wherein Q, T, X, and Z are independently selected from N or C, and wherein one of Q, T, X, and Z is N;

wherein L and M are independently selected from the group consisting of H, alkyl, alkoxy, aryl, substituted aryl, hydroxy, halogen, amino, alkylamino, nitro, cyano, CF₃, OCF₃, CONH₂, CONHR and NHCOR₁;

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₂-C₉ eyeloalkyl, substituted C₃-C₉ eyeloalkyl, polycyclic aliphatics,

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phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said method comprises steps:

reacting a compound with a formula:

with ammonium

hydroxide to form a compound with a formula:

reacting the compound with a formula:

with diammonium

sulfide to form a compound with a formula:

reacting the compound with a formula:

with a compound with

a formula:

to form a compound with a formula:

$$\begin{array}{c|c} O & T & Q & H & O & R \\ \hline & T & Q & H & O & M & R \\ R - NH & Z & NH_2 & M & H \end{array}$$
 to

cyclizing the compound with a formula: form to form a compound of Supragenus B.

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42. (Withdrawn – currently amended) A method of preparing a compound or salt thereof having the formula of Supreagenus B as defined in Claim 1 comprising:

$$\begin{array}{c|c} R_1 - H & T & Q & N & R_2 \\ \hline O & X & X & N & H & Supragenus B; \end{array}$$

wherein Q, T, X, and Z are independently selected from N or C, and wherein one of Q, T, X, and Z is N;

wherein L and M are independently selected from the group consisting of H, alkyl, alkoxy, aryl, substituted aryl, hydroxy, halogen, amino, alkylamino, nitro, cyano, CF₃, OCF₃, CONH₂, CONHR and NHCOR₁;

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R´ is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ eycloalkyl, substituted C₃-C₉ eycloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and

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substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said method comprises steps:

with ammonium

reacting a compound with a formula:

hydroxide to form a compound with a formula:

reacting the compound with a formula:

with diammonium

sulfide to form a compound with a formula:

reacting the compound with a formula:

with a compound with

a formula:

to form to form a compound of Supragenus B.

43. (Withdrawn - currently amended) A method of preparing a compound or salt thereof having the formula of Supragenus C as defined in Claim 1 comprising:

Supragenus C;

wherein Q, T, X, and Z are independently selected from N or C, and wherein one of Q, T, X, and Z is N;

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wherein L and M are independently selected from the group consisting of H, alkyl, alkoxy, aryl, substituted aryl, hydroxy, halogen, amino, alkylamino, nitro, cyano, CF₃, OCF₃, CONH₂, CONHR and NHCOR₁;

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁-and R₂-are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said method comprises steps:

reacting a compound with a formula: $O_2N \xrightarrow{T} O_2$ with ammonium hydroxide

$$O_2N \xrightarrow{X} Z NO_2$$
 NH_2

to form a compound with a formula:

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reacting the compound with a formula:

$$O_2N \xrightarrow{T} Q NO_2 NH_2$$

with diammonium

 O_2N Z NH_2 NH_2 NH_2

sulfide to form a compound with a formula:

 $O_2N \xrightarrow{T} Q NH_2$ $X \times Z NH_2$

reacting the compound with a formula:

with a compound with a

CI N.R

formula:

to form a compound with a formula:

$$O_2N_{\frac{1}{2}}^{\frac{1}{2}}$$
 $O_2N_{\frac{1}{2}}^{\frac{1}{2}}$
 $O_2N_{\frac{1}{2}}^{\frac{1}{2}}$

 $O_2N \xrightarrow{T} Q NH_2 O M$

cyclizing the compound with a formula:

with use

of an acid to form a compound with a formula:

 $O_2N \xrightarrow{T} Q N \xrightarrow{N} M \xrightarrow{N} R$

reducing the compound with a formula:

to form a

 $H_2N \xrightarrow{T} Q N M R$

compound with a formula:

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with an

reacting the compound with a formula:

acyl chloride to form a compound of Supragenus C.

44. (Withdrawn – currently amended) A method of preparing a compound or salt thereof having the formula of Supragenus C as defined in Claim 1 comprising:

$$\begin{array}{c|c} R_1 & T & Q & N & Q & N & R_2 \\ \hline & X & X & X & M & M & M & M \\ \hline & X & X & X & M & M & M & M \\ \hline \end{array}$$

Supragenus C;

wherein Q, T, X, and Z are independently selected from N or C, and wherein one of Q, T, X, and Z is N;

wherein L and M are independently selected from the group consisting of H, alkyl, alkoxy, aryl, substituted aryl, hydroxy, halogen, amino, alkylamino, nitro, cyano, CF₃, OCF₃, CONH₂, CONHR and NHCOR₁;

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl,

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dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ cycloalkyl, substituted C₃-C₉ cycloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said method comprises steps:

reacting a compound with a formula:

CI with ammonium hydroxide

$$O_2N \xrightarrow{T} Q NO_2 NH_2$$

to form a compound with a formula:

reacting the compound with a formula:

 $O_2N \xrightarrow{T} Q NO_2$ NH_2 with diammonium

$$O_2N$$
 X
 NH_2
 NH_2
 NH_2

sulfide to form a compound with a formula:

$$O_2N \xrightarrow{T} Q NH_2 NH_2 NH_2$$

reacting the compound with a formula:

with a compound with a

formula:

to form a compound with a formula:

$$O_2N \xrightarrow{T^{Q}} N \xrightarrow{N} N \xrightarrow{N} R$$

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$$O_2N \xrightarrow{T^{Q}} N \xrightarrow{N} M \xrightarrow{N} R$$

reducing the compound with a formula:

to form a

with an

compound with a formula:

reacting the compound with a formula: acyl chloride to form a compound of Supragenus C.

45. (Withdrawn – currently amended) A method of preparing a compound or salt thereof having the formula of Supragenus D as defined in Claim 1 compriring:

$$\begin{array}{c|c} R_1 & T & Q & N & H & R_2 \\ \hline HN & X & Z & N & M & O \end{array}$$

Supragenus D;

wherein Q, T, X, and Z are independently selected from N or C, and wherein one of Q, T, X, and Z is N;

wherein L and M are independently selected from the group consisting of H, alkyl, alkoxy, aryl, substituted aryl, hydroxy, halogen, amino, alkylamino, nitro, cyano, CF₃, OCF₃, CONH₂, CONHR and NHCOR₁;

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ eycloalkyl, substituted C₃-C₉ eycloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said

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substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH3, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉-cycloalkyl, substituted C₃-C₉-cycloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur:

wherein said method comprises steps:

 O_2N Z CI

reacting a compound with a formula:

with ammonium hydroxide

to form a compound with a formula:

reacting the compound with a formula:

with diammonium

sulfide to form a compound with a formula:

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reacting the compound with a formula:

rith a compound with a

form compound with formula: to

formula:

cyclizing the compound with a formula:

with use of

to form a

an acid to form a compound with a formula:

reducing the compound with a formula:

compound with a formula:

; and

reacting the compound with a formula: with an acyl

chloride to form a compound of Supragenus D.

46. (Withdrawn - currently amended) A method of preparing a compound or salt thereof having the formula of Supragenus D as defined in Claim 1 comprising:

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wherein Q, T, X, and Z are independently selected from N or C, and wherein one of Q, T, X, and Z is N;

wherein L and M are independently selected from the group consisting of H, alkyl, alkoxy, aryl, substituted aryl, hydroxy, halogen, amino, alkylamino, nitro, cyano, CF₃, OCF₃, CONH₂, CONHR and NHCOR₁;

wherein R is selected from the group consisting of H, C₁-C₅ alkyl, benzyl, p-fluorobenzyl and di-alkylamino alkyl, wherein said C₁-C₅ alkyl is selected from the group consisting of a straight chain, branched or cyclic alkyl;

wherein R₁ and R₂ are independently selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ eyeloalkyl, substituted C₂-C₉ eyeloalkyl, polycyclic aliphatic groups, substituted polycyclic aliphatic groups, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

wherein said substituted polycyclic aliphatic groups, substituted phenyl, substituted naphthyl and substituted heteroaryl contain 1-3 substituents, wherein said substituent is selected from the group consisting of H, halogens, polyhalogens, alkoxy group, substituted alkoxy, alkyl, substituted alkyl, dialkylaminoalkyl, hydroxyalkyl, carbonyl, OH, OCH₃, COOH, COOR' COR', CN, CF₃, OCF₃, NO₂, NR'R', NHCOR' and CONR'R'; and

wherein R' is selected from the group consisting of H, alkyl, substituted alkyl, C₃-C₉ eyeloalkyl, substituted C₃-C₉ eyeloalkyl, polycyclic aliphatics, phenyl, substituted phenyl, naphthyl, substituted naphthyl, heteroaryl and substituted heteroaryl, wherein said heteroaryl and said substituted heteroaryl contain 1-3 heteroatoms, wherein said heteroatom is independently selected from the group consisting of nitrogen, oxygen and sulfur;

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wherein said method comprises steps:

reacting a compound with a formula:

with ammonium hydroxide

 O_2N X Z NH_2

to form a compound with a formula:

reacting the compound with a formula:

with diammonium

Sulfide to form a compound with a formula: $O_2N \xrightarrow{T} O_2N \xrightarrow{X} Z$ NH_2

 $O_2N \xrightarrow{T} Q NH_2 NH_2$

reacting the compound with a formula:

with a compound with a

with an acyl

H NO

formula:

to form a compound with a formula:

 $O_2N \xrightarrow{T}^Q N \xrightarrow{N}_{H} NO_2$

reducing the compound with a formula:

compound with a formula:

reacting the compound with a formula:

chloride to form a compound of Supragenus D.

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47. (New) A pharmaceutical composition for treating or preventing an allergic reaction associated with increased IgE levels or inhibiting cellular proliferation in a mammal comprising any one or more of the compounds or salts thereof of Claim 1.

- 48. (New) The pharmaceutical composition of Claim 47, further comprising at least one additional ingredient which is active in reducing at least one symptom associated with said allergic reaction and/or cell proliferation.
 - 49. (New) A compound or salt thereof represented by the following formula: